

FORM PTO-1390
(REV. 5-93)U.S. DEPARTMENT OF COMMERCE
PATENT AND TRADEMARK OFFICE

ATTORNEY'S DOCKET NUMBER

KAS-157

U.S. APPLICATION NO. (if known, see 37 CFR 1.5)

09/936918

TRANSMITTAL LETTER TO THE UNITED STATES
DESIGNATED/ELECTED OFFICE (DO/EO/US)

International Application No.

PCT/JP00/01280

International Filing Date

March 3, 2000

Priority Date Claimed

Title of Invention AUTOMATIC ANALYZING APPARATUS

Applicant(s) for DO/EO/US S. NAKATA et al (see attached)

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a FIRST submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371.
3. ☒ This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).
4. ☒ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2))
 - a. ☐ is transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☒ has been transmitted by the International Bureau.
 - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☒ A translation of the International Application into English (35 U.S.C. 371(c)(2)).
7. ☐ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
 - a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☐ have been transmitted by the International Bureau.
 - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
 - d. ☐ have not been made and will not be made.
8. ☐ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☐ An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

Items 11. to 16. below concern other document(s) or information included:

11. ☒ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. ☐ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. ☐ A FIRST preliminary amendment.
☐ A SECOND or SUBSEQUENT preliminary amendment.
14. ☐ A substitute specification.
15. ☐ A change of power of attorney and/or address letter.
16. ☒ Other items or information:

☒ LIST OF INVENTORS' NAMES AND ADDRESSES.☒ THIS APPLICATION IS BEING FILED WITHOUT AN EXECUTED DECLARATION, WHICH WILL BE FILED LATER.

U.S. Application No. <i>if known, see 37 CFR 1.61</i>		International Application No.		Attorney's Docket Number	
09/936918		PCT/JP00/01280		KAS-157	

17. ☒ The following fees are submitted:

Basic National Fee (37 CFR 1.492 (a)(1)-(5)):

Search Report has been prepared by the EPO or JPO \$860.00

International preliminary examination fee paid to USPTO (37 CFR 1.482) \$690.00

No international preliminary examination fee (37 CFR 1.482) but international search fee paid to USPTO (37 CFR 1.445 (A)(2)) \$710.00

Neither international examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(A)(2)) paid to USPTO \$1000.00

International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(2) to (4) \$100.00

ENTER APPROPRIATE BASIC FEE AMOUNT		=	\$ 860.00		
------------------------------------	--	---	-----------	--	--

Surcharge of \$130.00 for furnishing the oath or declaration later than <input checked="" type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(e)).		+ \$	130.00		
--	--	------	--------	--	--

Claims	Number Filed	Number Extra	Rate		
Total	15 -20 =	0	x \$18.00	\$	0.00
Independent	2 - 3 =	0	x \$80.00	\$	0.00
Multiple dependent claim(s) (if applicable)				+ \$270.00	\$ 0.00
TOTAL OF ABOVE CALCULATIONS				=	\$ 990.00

Reduction by 1/2 for filing by small entity, if applicable. Verified Small Entity statement must also be filed. (Note 37 CFR 1.9, 1.27, 1.28).		\$	0.00		
SUBTOTAL		=	\$ 990.00		

Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f)).		+ \$	0.00		
TOTAL NATIONAL FEE		=	\$ 990.00		

Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property.		+ \$	0.00		
TOTAL FEES ENCLOSED		=	\$ 990.00		
				Amount to be:	
				Refunded \$	
				Charged \$	

a. ☒ A check in the amount of \$ 990.00 to cover the above fees is enclosed.

b. ☐ Please charge my Deposit Account No. 50-1417 in the amount of \$ _____ to cover the above fees. A duplicate copy of this sheet is enclosed.

c. ☒ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. 50-1417. A duplicate copy of this sheet is enclosed.

Note: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.


SEND ALL CORRESPONDENCE TO:

MATTINGLY, STANGER & MALUR, P.C.

1800 Diagonal Rd., Suite 370

Alexandria, Virginia 22314


(703) 684 -1120



Signature

John R. Mattingly

Name



24956

PATENT TRADEMARK OFFICE

30,293

Registration Number

FP242PCT-01
KAS-157

LIST OF INVENTORS' NAMES AND ADDRESSES

Shinpei NAKATA, Tokyo, JAPAN;

Takahiko YOTSUGI, Tokyo, JAPAN;

Hiroaki ISHIZAWA, Tokyo, JAPAN.

FP242PCT-01
KAS-157

United States National Phase Patent Application

Title of the Invention

ANALYSIS INFORMATION MANAGEMENT METHOD
AND ANALYSIS INFORMATION MANAGEMENT SYSTEM

Inventors

Shinpei NAKATA,
Takahiko YOTSUGI,
Hiroaki ISHIZAWA.

5/pjs

DESCRIPTION

ANALYSIS INFORMATION MANAGEMENT METHOD AND ANALYSIS
INFORMATION MANAGEMENT SYSTEM

FIELD OF THE INVENTION

The present invention relates to an analysis information management method and an analysis information management system, and more particularly, to an analysis information management method and an analysis information management system which are suitable for automatically setting analysis parameters required for conducting an analysis into an analyzer.

BACKGROUND ART

An automatically analyzing apparatus relies on a chemical reaction of a sample and a reagent introduced into an reaction container to quantitatively measure contents of the sample. For using the automatically analyzing apparatus, analysis parameters must be set into the automatically analyzing apparatus as analysis conditions, requiring a large number of steps for this work. Here, the analysis parameters refer to parameters for each testing item in the automatically analyzing apparatus, and basically include information such as the amount of dispensed sample, reagents used for an analysis and amounts thereof, the wavelength of absorbance under measurement, the type of reaction process and a method of calculating a

concentration, a standard sample and known concentration for use in a calibration, and the like. The information relates to reagents, and differs from one another depending on reagent makers dedicated to the manufacturing even for reagents used for analyzing the same testing item. In addition, even with reagents manufactured by the same reagent maker, contents of analysis parameters may result in different information for different manufacturing lots. Thus, analysis parameters to be set must be always the latest analysis parameters related to reagents set in the automatically analyzing apparatus.

As a first method of setting analysis parameters into an automatically analyzing apparatus, for example, as described in JP-A-11 211731, it is known that user sets parameters through a keyboard or the like as initial settings. Also, as a second method, for example, as described in JP-A-5-288756, it is known that reagent identification bar codes adhered on reagent containers are read by a bar code reader, and a variety of parameters are read from a computer based on identification codes inscribed in the bar codes to set the parameters into an automatically analyzing apparatus.

DISCLOSURE OF THE INVENTION

Here, the parameter setting method described in JP-A-11-211731 implies problems that a large number of input steps are required since parameters are manually set from a keyboard, and that correct parameter settings may not be

made in some cases due to erroneous inputs and the like. Particularly, in recent automatically analyzing apparatuses, the number of parameters and the number of screens for setting the parameters tend to increase due to the advancement of their functions, so that special knowledge is required for setting them without errors. Actually, dedicated persons in a reagent supplier or the like may occasionally set parameters, conduct subsequent calibration and measurements of accuracy management samples for confirming the operation before the parameters are provided to users. In this event, the reagent suppliers must set parameters for all automatically analyzing apparatus in all testing facilities to which their reagents are supplied. Therefore, the number of work steps is increased with an increase in a plurality of automatically analyzing apparatuses or a plurality of testing facilities, and corresponding automatically analyzing apparatuses.

On the other hand, the parameter setting method described in JP-A-5-288756 reads parameter information from an external computer based on bar codes adhered on reagent containers for automatic settings, wherein, however, a problem arises that bar codes must be adhered on reagent containers. In addition, after setting analysis parameters, the user is required to have special knowledge for determining whether or not the settings have been correctly made.

It is an object of the present invention to provide an analysis information management method and an

analysis information management system which are capable of readily setting analysis parameters for testing items subjected to an analysis using a reagent.

To achieve the above object, in a service center having a database connected to a plurality of automatically analyzing apparatus used in a plurality of facilities through communication lines for storing analysis parameters related to a plurality of reagents for use in the plurality of automatically analyzing apparatus used in the plurality of facilities, the present invention transfers analysis parameters for a testing item to be analyzed using a reagent to an automatically analyzing apparatus through the communication line in response to a request from the automatically analyzing apparatus. This method can readily set required analysis parameters even if a reagent container does not have means such as a bar code which allows identification of a reagent contained therein.

Also, to achieve the above object, the present invention comprises a communication unit for transmitting and receiving to and from a plurality of automatically analyzing apparatuses information such as information on analyses performed by the respective automatically analyzing apparatuses, a database for storing information such as the analysis information, an analysis information parsing unit for evaluating and parsing results analyzed by the automatically analyzing apparatuses using the analysis information stored in the database, and a reagent parameter registration unit for registering information on reagents in

the database, wherein the communication unit retrieves information on analysis parameters related to a managed reagent from the database in response to a request from the automatically analyzing apparatus, and transfers the information to the automatically analyzing apparatus. This configuration can readily set required analysis parameters even if a reagent container does not have means such as a bar code which allows identification of a reagent contained therein.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 is a block diagram illustrating the configuration of an analysis information management system using an analysis information management method according to one embodiment of the present invention;

Fig. 2 is an apparatus configuration diagram illustrating the configuration of an automatically analyzing apparatus for use in the analysis information management system according to one embodiment of the present invention;

Fig. 3 is a flow chart illustrating the contents of an analysis parameter setting process according to one embodiment of the present invention;

Fig. 4 is an explanatory diagram of a reagent information screen in the analysis parameter setting process according to one embodiment of the present invention; and

Fig. 5 is an explanatory diagram of a reagent listing screen in the analysis parameter setting process

according to one embodiment of the present invention.

BEST MODE FOR CARRYING OUT THE INVENTION

In the following, an analysis information management method and an analysis information management system according to one embodiment of the present invention will be described with reference to Figs. 1 to 5.

First, the configuration of a system for use with the analysis information management method according to this embodiment will be described with reference to Fig. 1.

Fig. 1 is a block diagram illustrating the configuration of an analysis information management system which uses the analysis information management method according to one embodiment of the present invention.

A service center 10 is connected to all automatically analyzing apparatuses 100A, 100B, ..., 100Z in testing facilities, which have concluded a service agreement, through communication lines L. In each of the testing facilities, one or a plurality of automatically analyzing apparatuses 100 are installed such that the automatically analyzing apparatuses 100A, 100B are installed in a testing facility 1, and the automatically analyzing apparatus 100Z is installed in a testing facility M. The service center 10 is also connected to a plurality of reagent suppliers 20A, 20B, ..., 20N through the communication lines L.

The service center 10 intensively manages analysis information such as analysis parameters related to reagents,

results of analyses made by the respective automatically analyzing apparatuses, and the like. The service center 10 comprises a communication unit 12, an analysis information parsing unit 14, a database 16, and a reagent parameter registration unit 18.

The communication unit 12 transmits and receives analysis information between the automatically analyzing apparatuses 100 connected thereto through the communication lines L. The analysis information parsing unit 14 evaluates and parses analysis information such as results of analyses sent thereto from the connected automatically analyzing apparatuses 100. The database 16 is connected to the automatically analyzing apparatuses 100 through the communication lines L for unitarily managing all reagents available in all the associated automatically analyzing apparatuses 100 and analysis parameters related to the reagents, and analysis information on the associated automatically analyzing apparatuses 100, for example, information such as reagents used in analyses and analysis parameters, results of calibrations, results of analyses on accuracy management samples, and the like. The reagent parameter registration unit 18 is a unit for registering information related to reagents such as reagent information comprised of the names of testing items under analysis using reagents, reagent lot Nos, and the like, analysis parameters related to the reagents, and automatically analyzing apparatuses available with the reagents, and the like in the database 16.

The reagent suppliers 20 are dealers which supply their reagents to the respective facilities. The reagent suppliers 20 have concluded an agreement with the service center 10, in which the reagent suppliers 20 register the database 16 of the service center 10 through the reagent parameter registration unit 18 when their reagents or analysis parameters related to the reagents need be added or updated, within those available to all the automatically analyzing apparatuses 100 administered by the service center 10, due to the development of new reagents capable of analyzing new testing items, and a change in manufacturing lots of reagents even if they are used for existing testing items. In this way, the service center 10 holds the most recent information related to reagents.

In the automatically analyzing apparatus 100, analysis parameters must have been set for testing items which are to be analyzed using reagents set in the automatically analyzing apparatus 100 before starting the analysis. For setting a novel reagent supplied newly from a reagent supplier 20 into the automatically analyzing apparatus 100, associated analysis parameters may not have been set in the automatically analyzing apparatus 100. In this event, the automatically analyzing apparatus 100 performs an operation for requesting the service center 10 to send through the communication line L a list on which one can view all reagents available for the automatically analyzing apparatus 100. The service center 10 searches the database 16 to create a list of reagents available in the

automatically analyzing apparatus 100, which has made the request, from among the database 16, and transmits the list to the automatically analyzing apparatus 100 through the communication line L. The automatically analyzing apparatus 100 displays the received reagent list. As the user selects a reagent set in the automatically analyzing apparatus 100 from the list, the automatically analyzing apparatus 100 transmits the selected information to the service center 10. Upon receipt of the selected information, the service center 10 searches the database 16 for associated analysis parameters which are sent to the automatically analyzing apparatus 100 through the communication line L. The automatically analyzing apparatus 100 sets the received analyzing parameters.

While in the foregoing example, the reagent list is transmitted to the automatically analyzing apparatus 100, the service center 10 may have a home page for registering the reagent list such that the list can be viewed from the automatically analyzing apparatus 100.

Next, the configuration of an automatically analyzing apparatus for use in the analysis information management system according to this embodiment will be described with reference to Fig. 2.

Fig. 2 is an apparatus configuration diagram illustrating the configuration of the automatically analyzing apparatus for use in the analysis information management system according to one embodiment of the present invention.

A reaction container 112 is provided on the outer periphery of a reaction disk 110. The entire reaction disk 110 is held at a predetermined temperature by an incubation bath 114. A large number of sample cups 122 are disposed in a sample disk unit 120. A sample in a sample cup 122 is extracted as required through a nozzle 128 of a sample pipetting unit 126 which is controlled by a sample dispensing unit 124, and injected into the reaction container 112 at a sample dispensing position.

Reagent bottles with bar code labels are disposed in reagent disk units 130A, 130B. A bar code reader 132A, 132B is associated with each of the reagent disk units 130A, 130B for reading a bar code upon registration of a reagent to register reagent bottle information corresponding to a position. The reagent bottle information is comprised of information such as the type of reagent, lot No., expiration date, remaining reagent amount, and the like.

Each of the reagent disks is also provided with a second reagent pipetting unit 142A and a first reagent pipetting unit 142B controlled by the sample dispensing unit 140. The second reagent pipetting unit 142A pipets a reagent from the reagent disk using a nozzle 144A, and delivers it to the reaction container 112. The first reagent pipetting unit 142B in turn pipets a reagent from the reagent disk using a nozzle 144B and delivers it to the reaction container 112. Stirrers 146A, 146B are disposed in close proximity to the reagent disk units 130A, 130B for stirring a sample and a reagent within the reaction

container 112. The reaction container 112 which contains an object under photometry is disposed between an analysis unit comprised of a multi-wavelength photometer 150 and a light source 152. A washing unit 158 washes the reaction container 112 with a washing solution supplied from a washing water pump 156.

Further, as a control system and a signal processing system, the automatically analyzing apparatus 100 comprises a microcomputer 160, an interface 162, a Log (logarithm) converter 164, and an A/D converter 166. Also, the automatically analyzing apparatus 100 comprises a printer 170 for printing, a CRT 172 for display, a hard disk 174 as a storage device, and a manipulation panel for inputting (a keyboard or a pointing device such as a touch screen, a mouse, and the like) 176. The automatically analyzing apparatus 100 further comprises a communication unit 180 for communicating with the service center 10 through the communication line L.

Next, description will be made on the operation of the automatically analyzing apparatus 100 illustrated in Fig. 2.

A sample put in a sample cup 122 is dispensed to the reaction container 112 by a predetermined amount using the nozzle 128 of the sample pipetting unit 126 in accordance with analysis parameters stored in a memory of the microcomputer 160.

Next, the reaction container 112 dispensed with the sample is transported to a reagent dispensing position

by rotating the reaction disk 110. Subsequently, reagents are inputted through the manipulation panel 176, and are dispensed by predetermined amounts to the reaction container 112, dispensed with the sample, using the nozzles 144A, 144B of the reagent pipetting units 142A and 142B in accordance with analysis parameters stored in the microcomputer 160. Subsequently, the sample and reagents are stirred by the stirrers 146A, 146B for mixing.

When the reaction container 112 traverses a photometric position, the multi-wavelength photometer 150 measures the absorption. The measured absorption is captured into the microcomputer 160 through the Log converter 164, A/D converter 166 and interface 162. The absorption is converted to concentration data based on a calibration curve created from the absorption of a standard sample solution which has been previously measured by an analysis method specified for each item. The measured component concentration data is outputted to the printer 170 and to a screen of the CRT 172. The result of measurement may also be transmitted to the outside through the communication unit 180.

In the foregoing measurement principles, the user performs on the screen (CRT) 172 settings of various parameters required for the measurement, registration of the sample, and confirmation of the result of analysis.

Reagent bottles may or may not have a bar code adhered on the side surface thereof. A reagent bar code describes reagent bottle identification information, reagent

lot No., expiration date, and the like. As illustrated in Fig. 2, when the reagent bar code readers 132A, 132B are provided, bar codes are read from reagent bottles at all positions on the reagent disks, for example, when the reagent disks 130A, 130B are closed with lids, to register reagent bottle information corresponding to each of the positions. On the other hand, when bar codes are not adhered on bottles, reagent bottle information is registered by the user who manually enters information included in bar codes from the manipulation panel 176. The reagent bottle information is comprised of information such as the type of reagent, lot No., expiration date, remaining amount of reagent, and the like. Testing items available for analysis are determined by reagents set in the reagent disks 130A, 130B, and analysis parameters are required for defining analysis conditions therefor. Also, other methods of setting analysis parameters may involve specifying testing items and manually entering required parameters from the manipulation panel 176, reading parameters from FD or bar sheet, and the like.

Next, the contents of analysis parameter setting process in the analysis information management method according to this embodiment will be described with reference to Figs. 3 to 5.

First, the contents of the analysis parameter setting process according to this embodiment will be described with reference to Fig. 3.

Fig. 3 is a flow chart illustrating the contents

of the analysis parameter setting process according to one embodiment of the present invention.

Here, description will be made on the contents of the process from setting of reagent bottles on the reagent disks 130A, 130B of the automatically analyzing apparatus 100 to setting of analysis parameters associated therewith.

At step s200, the operator sets reagent bottles on the reagent disks 130A, 130B of the automatically analyzing apparatus 100, and for example, closes the lids of the reagent disks 130A, 130B, causing the microcomputer 160 of the automatic analysis apparatus 100 to read bar codes on the reagent bottles at all positions on the reagent disks 130A, 130B from the bar code readers 132A, 132B.

Then, at step s205, the microcomputer 160 searches stored reagent bottle information and analysis parameter information from the read bar code information.

Assuming herein that a reagent screen is opened at the automatically analyzing apparatus 100, the CRT 172 displays on a reagent information screen 172A information related to the reagent set at each of the positions on the reagent disks, for example, as illustrated in Fig. 4.

Here, referring to Fig. 4, an example of reagent information screen will be described in this embodiment.

Fig. 4 is an explanatory diagram of the reagent information screen in the analysis parameter setting process according to one embodiment of the present invention.

The reagent information screen 172A is comprised of Position, Testing Item, Reagent Type, Reagent Lot No.,

Reagent Bottle No., Presence/Absence of Analysis Parameters, Reagent Information Serial #, and the like. Position indicates the position at which a reagent bottle is set on the reagent disk units 130A, 130B, and gives a serial number such as No. 1, No. 2, and the like, by way of example. The shown example indicates a reagent set at Position 1 has a testing item name "ALP," reagent type "R1," reagent lot No. "100001," reagent bottle No. "10001," and analysis parameters "presence." The reagent information serial # will be described later with reference to Fig. 5. For example, for a reagent at Position "3," the table shows that analysis parameters are "absent." Also, at Position "4," the table does not at all display information such as Position, Testing Item, Reagent Type, Reagent Lot No., Reagent Bottle No., Presence/Absence of Analysis Parameters, and the like, meaning that no bar code is adhered on a reagent bottle, so that the information is displayed as blank.

When no bar code is adhered on a reagent bottle, for example, as Position 4, the operator sees the reagent bottle and enters information which allows identification of the reagent such as the reagent lot No. or the like in blanked fields on the reagent information screen 172A.

At step s210, the microcomputer 160 determines whether or not reagent bar code information has been manually entered on the reagent screen. The process continues to step s215 when entered, and to step s220 when not entered.

As described above, when information is manually entered for the reagent at Position "4," the microcomputer 160 searches stored reagent bottle information for information on the reagent, and displays the information, if found, on the reagent information screen 172A at step s215. On the other hand, if analysis parameters have been stored for a testing item which is analyzed with the reagent, "presence" is displayed in the field of Presence/Absence of Analysis Parameters. When a novel reagent is set and therefore no associated analysis parameters have been stored, the absence of analysis parameters is displayed as Position 3, in which case analysis parameters must be newly set before an analysis is started. In this event, the operator depresses a "reagent list download button" positioned in a lower right region in the reagent information screen 172A illustrated in Fig. 4.

At step s220, the microcomputer 160 determines whether or not the operator has instructed to download a reagent list. The process continues to step s225 when instructed, and to s240 when not instructed. As described above, as the operator depresses the "reagent list download button," the process continues to step s225 assuming that the operator has instructed to download the reagent list.

At step s225, the microcomputer 160 transmits a reagent list download request to the service center 10 through the communication line L. In this event, the microcomputer 160 also transmits identification information of the automatically analyzing apparatus 100 such as the

model number to the service center 10.

The service center 10 searches for all reagents available in the automatically analyzing apparatus 100, which has made the reagent list download request, collects them in the form of list, and transmits the list to the automatically analyzing apparatus 100 through the communication line L.

At step s230, the microcomputer 160 of the automatically analyzing apparatus 100 is monitoring until the reagent list has been downloaded, and upon completion of the downloading, the microcomputer 160 displays a reagent list as in Fig. 5 on the CRT 172 as a reagent list screen 172B at step s235.

Now, an example of the reagent list screen in this embodiment will be described with reference to Fig. 5.

Fig. 5 is an explanatory diagram of the reagent list screen in the analysis parameter setting process according to one embodiment of the present invention.

The reagent list screen 172B is comprised of Reagent Information Serial #, Reagent Maker Name, Testing Item, Reagent Lot No., and the like. Reagent Information Serial # is a serial number for unitarily managing reagents. The shown example indicates that a reagent having Reagent Information Serial # "20001" has Reagent Maker Name "AAA," Testing Item Name "AST," and Reagent Lot No. "00001." The operator views this reagent list to search for a reagent for which analysis parameters must be newly set, and enters a testing item name and a reagent information serial number in

fields at corresponding positions on the reagent screen in Fig. 3. For example, when a reagent at Position "4" in Fig. 4 is manufactured by a reagent maker named "AAA" and used for a testing item labeled "AST," and has Reagent Lot No. "00001," the operator enters "AST" in Testing Item Name, and "2002" in Reagent Information Serial # on Position "4" in Fig. 4. This causes information such as Reagent Type, Reagent Lot No. and the like to be entered automatically. This processing is performed for all reagents for which analysis parameters have not been stored and therefore must be newly set. Upon completion of this processing, the operator depresses a "parameter download button" in a lower right region on the reagent information screen 172A in Fig. 4.

At step s240, the microcomputer 160 of the automatically analyzing apparatus 100 determines whether or not the operator has selected reagent information, and whether or not the operator has instructed to download analysis parameters. The process continues to step s245 when the operator has selected or instructed, and to step s260 when not.

When the operator has selected reagent information or instructed to download analysis parameters, the microcomputer 160 creates an analysis parameter download request which is transmitted to the service center 10 through the communication line L at step s245. The service center 10 searches for all analysis parameters requested for downloading and transmits the analysis parameters to the

automatically analyzing apparatus 100.

At step s250, the microcomputer 160 of the automatically analyzing apparatus 100 is monitoring until the requested analysis parameters have been downloaded. Upon completion of the downloading, the microcomputer 160 stores the received analysis parameters at step s255. As the analysis parameters have been stored, "presence" is displayed in the field "Presence/Absence of Analysis Parameters" at the associated position on the reagent information screen 172A shown in Fig. 4.

At step s260, the microcomputer 160 determines whether or not an analysis start key has been depressed. As the analysis start key is depressed, analysis parameters are stored in the memory of the microcomputer 160 upon start of the analysis for conducting the analysis based on the analysis parameters at step s265.

In the foregoing description, when a reagent bar code is adhered on a reagent bottle, the reagent screen is opened after the reagent bar code has been read, resulting in a search made for reagent bottle information and analysis parameters based on the bar code information. If information has been previously stored, the information is displayed as Position 1 or 2 in Fig. 4. On the other hand, if a novel reagent is set on the reagent disk, reagent bottle information is stored. When associated analysis parameters are present, the reagent bottle information is displayed on the screen in a similar manner. If associated analysis parameters have not been stored, "absence" of

analysis parameters is displayed as Position 3 in Fig. 4. Even in this event, analysis parameters can be downloaded and set as is the case with a reagent bottle without a bar code adhered thereon.

If a reagent bar code includes information such as the name of the reagent maker, testing item name, and the like which allows identification of analysis parameters, a parameter download request may be automatically made for the reagent, such that analysis parameters can be set without requiring the user to download a reagent list to select associated reagent information.

Next, a method of verifying that analysis parameters have been correctly set in the automatically analyzing apparatus will be described with reference to Fig. 1.

The automatically analyzing apparatus 100 conducts an analysis based on the analysis parameters set at step s265 in Fig. 3. When a new reagent has been introduced and analysis parameters associated therewith have been newly set, the operator actually performs a calibration to create a calibration curve before an accuracy management sample is analyzed. The controller 160 of the automatically analyzing apparatus 100 transmits the result of calibration and the result of analysis on the accuracy management sample, as well as analysis information such as reagents used therein and analysis parameters to the service center 10 through the communication line L.

The service center 10, upon receipt of the analysis

information from each automatically analyzing apparatus 100 through the communication unit 12, classifies the received analysis information by testing facilities or automatically analyzing apparatuses, and saves the classified analysis information in the database 16 for management. The analysis information parsing unit 14 of the service center 10 retrieves the results of analyses on accuracy management samples measured using the same reagent, managed in the database 16, and calculates an average value which is defined as a standard value. Then, upon determination that the currently received result of analysis on the accuracy management sample has been acquired using newly set analysis parameters, the analysis information parsing unit 14 calculates a deviation between the previously found standard value and the currently received analysis result, and determines whether or not the deviation is within an allowable range. If the deviation is out of the allowable range, the analysis information parsing unit 14 confirms, for example, whether correct analysis parameters are used for reagents used in the analysis, whether the set parameters are inconsistent in the contents, and the like. Then, the analysis information parsing unit 14 creates a report which summarizes the result of verification, countermeasures to be taken by the user, and the like, and transmits the report to the automatically analyzing apparatus 100.

The operator of the automatically analyzing apparatus, who has received the report, relies on the

contents of the report to, for example, start a testing work if no defect is found in the result of verification, and takes action in accordance with the contents of the report if any defect is found.

In the foregoing manner, the analysis information such as information on the results of measurements made by the automatically analyzing apparatus is transmitted to the service center, such that the results are parsed and evaluated in the service center, thereby making it possible to verify the operation of the apparatus from a global view point and to support the user of the automatically analyzing apparatus who does not have special knowledge to readily verify that analysis parameters have been correctly set.

Each time the result of analysis on the accuracy management sample is received, the analysis information parsing unit 14 calculates a standard value and a deviation, and transmits the result of calculation to the automatically analyzing apparatus 100 which has transmitted the result of analysis in the form of report in a similar manner if any defect is found. On the other hand, if no defect is found, the analysis information parsing unit 14 stores the result of calculation in the database 16, and periodically creates a report which is transmitted to the automatically analyzing apparatus 100.

In the foregoing description, the analysis information parsing unit 14 conducts up to the calculation of a deviation between the result of measurement by the automatically analyzing apparatus 100 and the standard value

in the entirety, and sends the result of calculation to the automatically analyzing apparatus 100 as a report. Alternatively, the analysis information parsing unit 14 may periodically calculate standard values and the like of all automatically analyzing apparatuses administrated thereby, and periodically send the information to the automatically analyzing apparatuses 100 as technical information. In this way, the operator of the automatically analyzing apparatus 100 can evaluate, for example, the result of measurement of an accuracy management sample based on the technical information, and can verify by himself that newly set analysis parameters are being correctly used.

In the foregoing description, while the database 16 of the service center 10 is configured to manage information on analyses such as reagents, analysis parameters, results of measurements, and the like, the database 16 may store, by version, programs for controlling all automatically analyzing apparatuses administered thereby, in addition to the foregoing information. In this way, for example, as a new reagent is introduced, a specified analysis method is added for each testing item, so that if an addition, update and the like are required for a program for converting to concentration data based on the analysis method, the program can be downloaded from the service center for automatic installation into the automatically analyzing apparatus, as is the case with the analysis parameters. Further, since the database 16 of the service center 10 can also manage information on versions of

programs installed in the automatically analyzing apparatuses administered thereby, it is possible to correctly keep track of the current states of the administered automatically analyzing apparatuses and to support a change in programs as mentioned in a timely manner.

As described above, according to this embodiment, even under condition that there are a plurality of facilities having a plurality of automatically analyzing apparatuses using reagents from a plurality of reagent suppliers, even if reagent containers do not have bar codes adhered thereon for allowing identification of reagents therein, it is possible to readily set analysis parameters for testing items to be analyzed using the reagents, and to readily verify that the analysis parameters have been correctly set, thereby improving the efficiency and reliability for the testing work conducted by the automatically analyzing apparatus. Also, when viewed from the reagent suppliers, it is expected to improve the efficiency in works related to the distribution of analysis parameters to testing facilities, and the like.

INDUSTRIAL AVAILABILITY

According to the present invention, analysis parameters can be readily set for testing items which are analyzed using reagents.

CLAIMS

1. An analysis information management method using a service center having a database (16) connected to a plurality of automatically analyzing apparatuses (100) used in a plurality of facilities through communication lines for storing analysis parameters related to a plurality of reagents for use in the plurality of automatically analyzing apparatus used in the plurality of facilities,

characterized by transferring analysis parameters for a testing item to be analyzed using a reagent to an automatically analyzing apparatus through the communication line in response to a request from said automatically analyzing apparatus.

2. An analysis information management method according to claim 1, characterized in that:

said automatically analyzing apparatus (100) automatically sets the transferred analysis parameters.

3. An analysis information management method according to claim 1, characterized in that:

said database (16) stores analysis parameters related to reagents from a plurality of reagent suppliers.

4. An analysis information management method according to claim 1, characterized in that:

said service center (10), responsive to a request

from said automatically analyzing apparatus, creates a list of reagents available in said requesting automatically analyzing apparatus within information on reagents stored in said database, and supplies said automatically analyzing apparatus with the list through said communication line.

5. An analysis information management method according to claim 4, characterized in that:

said service center (10), responsive to a selection of an associated reagent from said list made by a user of said automatically analyzing apparatus, transfers analysis parameters for a testing item to be analyzed using the selected reagent to said automatically analyzing apparatus through said communication line.

6. An analysis information management method according to claim 1, characterized in that:

when a reagent supplier (20) supplies a novel reagent or a reagent in a new lot to a user of an automatically analyzing apparatus, said reagent supplier registers said database with information related to said reagent, such as said reagent, automatically analyzing apparatuses capable of using said reagent, and analysis parameters for said reagent prior to the supply.

7. An analysis information management method according to claim 1, characterized in that:

said service center (10) classifies and stores

information on analyses such as results of calibrations measured by automatically analyzing apparatuses, results of analyses on accuracy management samples, reagents used in analyses, and analysis parameters for each facility or for each automatically analyzing apparatus; and

calculates, based on the stored information on the results of analyses, a standard value for results of analyses on accuracy management samples using the same reagents in all automatically analyzing apparatuses in all facilities administered by said service center, so that when a certain automatically analyzing apparatus administered said service center has newly analyzed an accuracy management sample, said service center calculates a deviation between the result of analysis and said standard value for evaluation to verify that analysis parameters used in the analysis are correct.

8. An analysis information management method according to claim 7, characterized in that:

said service center (10), upon determination that the result of analysis on an accuracy management sample transferred thereto from an automatically analyzing apparatus was derived using newly set analysis parameters, summarizes the result of verification in a report, and transmits the report to the automatically analyzing apparatus through the communication line.

9. An analysis information management method

according to claim 7, characterized in that:

each time said service center (10) receives the result of analysis on an accuracy management sample from an automatically analyzing apparatus, said service center calculates a deviation from said standard value, and transmits the result of analysis to the automatically analyzing apparatus through the communication line if any defect is recognized.

10. An analysis information management method according to claim 9, characterized in that:

when no defect is recognized in the result of analysis, said service center (10) stores the result of analysis, periodically creates a report, and transmits the report to the automatically analyzing apparatuses through the communication lines.

11. An analysis information management method according to claim 7, characterized in that:

said service center (10) periodically calculates said standard value, and transmits said standard value to the automatically analyzing apparatuses through the communication lines as technical information.

12. An analysis information management method according to claim 1, characterized in that:

said service center (10) stores and manages by version programs for controlling the automatically analyzing

apparatuses administered thereby, and automatically installs a program of a requested version in response to a request from an automatically analyzing apparatus administered thereby.

13. An analysis information management system comprising:

a communication unit (12) for transmitting and receiving among a plurality of automatically analyzing apparatuses information such as information on analyses performed by the respective automatically analyzing apparatuses;

a database (16) for storing information such as the analysis information;

an analysis information parsing unit (14) for evaluating and parsing results analyzed by said automatically analyzing apparatuses using the analysis information stored in said database; and

a reagent parameter registration unit (18) for registering information on reagents in said database,

characterized in that said communication unit (12) retrieves information on analysis parameters related to a managed reagent from said database in response to a request from said automatically analyzing apparatus, and transfers the information to said automatically analyzing apparatus.

14. An analysis information management system according to claim 13, characterized in that:

said communication unit (12) retrieves information on a list of managed reagent from said database in response to a request from said automatically analyzing apparatus, and transfers the information to said automatically analyzing apparatus.

15. An analysis information management system according to claim 13, characterized in that:

said analysis information parsing unit (14) calculates a standard value for results of analyses on accuracy management samples made using the same reagent in all automatically analyzing apparatuses in all facilities administered by said service center based on analysis information such as results of calibrations measured by the automatically analyzing apparatuses, results of analyses on the accuracy management samples, reagents used in the analyses, and analysis parameters, said analysis information stored in said database, so that when a certain automatically analyzing apparatus administered by said service center has newly analyzed an accuracy management sample, said analysis information parsing unit calculates a deviation between the result of analysis and said standard value for evaluation to verify that the analysis parameters used in the analysis are correct.

FIG. 1

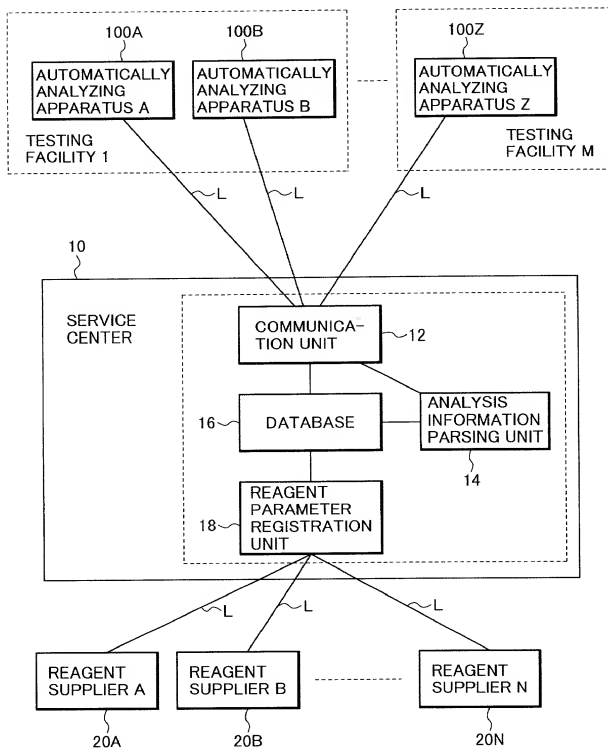


FIG. 2

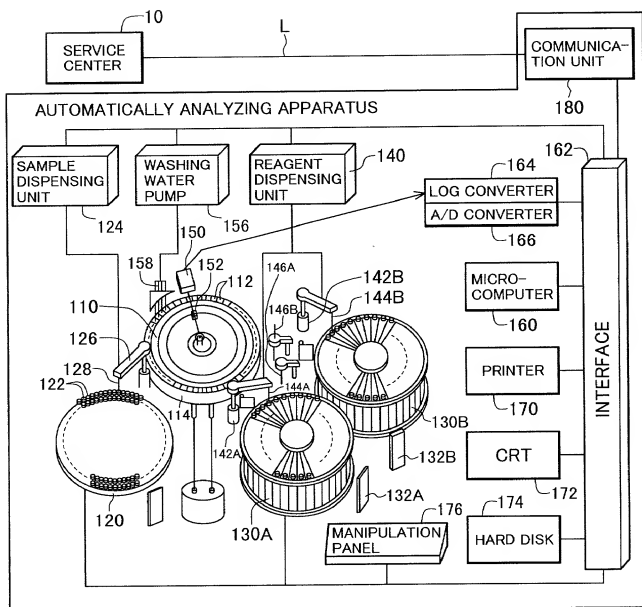
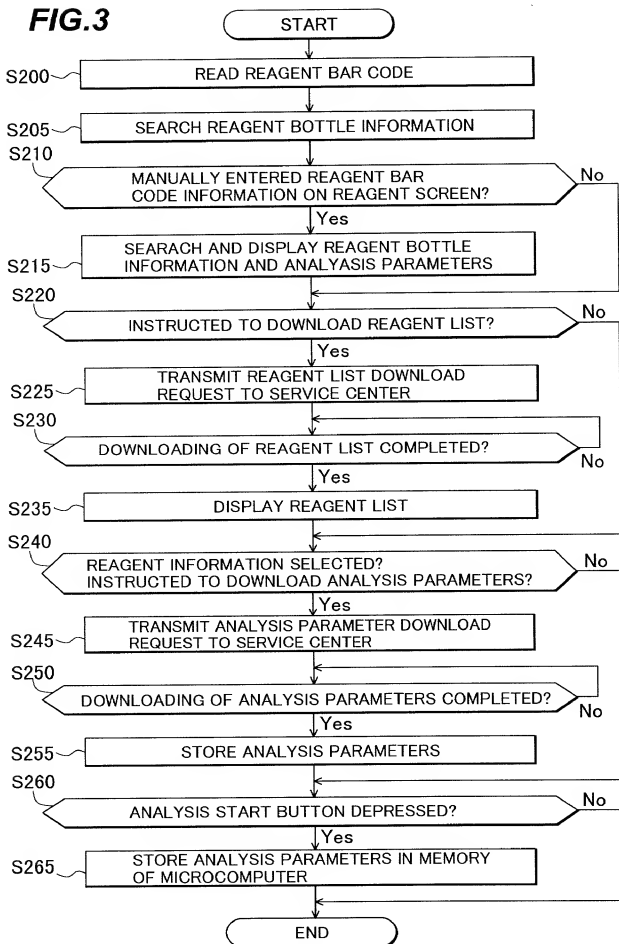


FIG.3

172A

POSITION	TESTING ITEM NAME	REAGENT TYPE	REAGENT LOT NO.	REAGENT BOTTLE NO.	PRESENCE/ ABSENCE OF ANALYSIS PARAMETERS	REAGENT INFORMATION SERIAL NO.
1	ALP	R1	100001	10001	○	
2	ALP	R2	100001	10002	○	
3		R1	100010	10010		
4						
:						

REAGENT LIST
DOWN LOAD

PARAMETER
DOWNLOAD

FIG.5

172B

REAGENT LIST SCREEN						
REAGENT INFORMATION SERIAL NO.	REAGENT MAKER NAME	TESTING ITEM NAME	REAGENT LOT NO.	...		
2001	AAA	AST	00001			
2002	AAA	AST	00002			
2003	BBB	AST	00010			
:						



FP282
(11905322)
#3

Under the Paperwork Reduction Act of 1995, no persons are required to provide information unless it displays a valid OMB control number.

PTO/SB/106(8-96)
Approved for use through 9/30/98. OMB 0651-0032

Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE

Declaration and Power of Attorney For Patent Application

特許出願宣言書及び委任状

Japanese Language Declaration

日本語宣言書

下記の氏名の発明者として、私は以下の通り宣言します。

As a below named inventor, I hereby declare that:

私の住所、私書箱、国籍は下記の私の氏名の後に記載された通りです。

My residence, post office address and citizenship are as stated next to my name.

下記の名称の発明に関して請求範囲に記載され、特許出願している発明内容について、私が最初かつ唯一の発明者（下記の氏名が一つの場合）もしくは最初かつ共同発明者であると（下記の名称が複数の場合）信じています。

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled

ANALYSIS INFORMATION MANAGEMENT METHOD AND

ANALYSIS INFORMATION MANAGEMENT SYSTEM

上記発明の明細書（下記の欄で×印がついていない場合は、本書に添付）は、

The specification of which is attached hereto unless the following box is checked:

☐ __月__日に提出され、米国出願番号または特許協定条約国際出願番号を____とし、
(該当する場合) _____に訂正されました。

☒ was filed on March 3, 2000
as United States Application Number or
PCT International Application Number
PCT/JPO00/01280 and was amended on
____ (if applicable).

私は、特許請求範囲を含む上記訂正後の明細書を検討し、内容を理解していることをここに表明します。

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

私は、連邦規則法典第37編第1条56項に定義されるとおり、特許資格の有無について重要な情報を開示する義務があることを認めます。

I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, Section 1.56.

Under the Paperwork Reduction Act of 1996, no persons are required to respond to a collection of information unless it displays a valid OMB control number.

Japanese Language Declaration
(日本語宣言書)

私は、米国法典第35編119条(a)-(d)項又は365条(b)項に基づき下記の、米国以外の国の少なくとも一カ国を指定している特許協力条約365(a)項に基づく国際出願、又は外国での特許出願もしくは発明者証の出願についての外国優先権をここに主張するとともに、優先権を主張している、本出願の前に出願された特許または発明者証の外国出願を以下に、枠内をマークすることで、示している。

Prior Foreign Application(s)

外国での先行出願

(Number) (番号)	(Country) (国名)
(Number) (番号)	(Country) (国名)

私は、第35編米国法典119条(e)項に基づいて下記の米国特許出願規定に記載された権利をここに主張いたします。

(Application No.) (出願番号)	(Filing Date) (出願日)
-----------------------------	------------------------

私は、下記の米国法典第35編120条に基づいて下記の米国特許出願に記載された権利、又は米国を指定している特許協力条約365条(c)項に基づき権利をここに主張します。また、本出願の各請求範囲の内容が米国法典第35編112条第1項又は特許協力条約で規定された方法で先行する米国特許出願に開示されていない限り、その先行米国出願書提出日以降で本出願書の日本国内または特許協力条約国際提出日までの期間中に入手された、連邦規則法典第37編1.56条で定義された特許資格の有無に関する重要な情報について開示義務があることを認識しています。

(Application No.) (出願番号)	(Filing Date) (出願日)
(Application No.) (出願番号)	(Filing Date) (出願日)

私は、私自身の知識に基づいて本宣言書中で私が行なう表明が真実であり、かつ私の入手した情報と私の信じることに基づき表明が全て真実であると信じていること、さらに故意になされた虚偽の表明及び同等の行為は米国法典第18編第1001条に基づき、罰金または拘禁、もしくはその両方により処罰されること、そしてそのような故意による虚偽の声明を行えば、出願した、又は既に許可された特許の有効性が失われることを認識し、よってここに上記のごとく宣誓を致します。

I hereby claim foreign priority under Title 35, United States Code, Section 119 (a)-(d) or 365(b) of any foreign application(s) for patent or inventor's certificate, or 365(a) of any PCT international application which designated at least one country other than the United States, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or PCT International application having a filing date before that of the application on which priority is claimed.

Priority Not Claimed

優先権主張なし

I hereby claim the benefit under Title 35, United States Code, Section 119(e) of any United States provisional application(s) listed below.

(Application No.) (出願番号)	(Filing Date) (出願日)
-----------------------------	------------------------

I hereby claim the benefit under Title 35, United States Code, Section 120 of any United States application(s), or 365(c) of any PCT international application designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application in the manner provided by the first paragraph of Title 35, United States Code Section 112, I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, Section 1.56 which became available between the filing date of the prior application and the national or PCT international filing date of application.

(Status: Patented, Pending, Abandoned) (現況: 特許許可済、係属中、放棄済)
(Status: Patented, Pending, Abandoned) (現況: 特許許可済、係属中、放棄済)

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it displays a valid OMB control number.

Japanese Language Declaration (日本語宣言書)

委任状： 私は下記の発明者として、本出願に関する一切の手続きを米特許商標局に対して遂行する弁理士または代理人として、下記の者を指名いたします。(弁理士、または代理人の氏名及び登録番号を明記のこと)

POWER OF ATTORNEY: As a named inventor, I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and transact all business in the Patent and Trademark Office connected therewith (list name and registration number)

John R. Mattingly, Reg. No. 30,293; Daniel J. Stanger, Reg. No. 32,846; Shrinath Malur, Reg. No. 34,663; Gene W. Stockman, Reg. No. 21,021; Jeffrey M. Ketchum, Reg. No. 31,174; and Scott W. Brickner, Reg. No. 34,553.

書類送付先

Send Correspondence to:

MATTINGLY, STANGER & MALUR, P.C.

1800 Diagonal Road, Suite 370

Alexandria, Virginia 22314

直接電話連絡先：(氏名及び電話番号)

Direct Telephone Calls to: (name and telephone number)

Telephone: (703) 684-1120

Fax: (703) 684-1157

唯一または第一発明者	1 - 80	Full name of sole or first inventor
発明者の署名	日付	Shinsai NAKATA
住所	Inventor's signature	Date
国籍	Shinsai Nakata	9/21/2001
私書箱	Residence	JPX
	Mito, Japan	
	Citizenship	
	Japan	
	Post Office Address	
	c/o Hitachi, Ltd., Intellectual Property Group	
	New Marunouchi Bldg. 5-1, Marunouchi 1-chome,	
	Chiyoda-ku, Tokyo 100-8220, Japan	

(第二以降の共同発明者についても同様に記載し、署名をすること)

(Supply similar information and signature for second and subsequent joint inventors.)

Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it displays a valid OMB control number.

第二共同発明者		2-00		Full name of second joint inventor, if any Takehiko YOTSUGI	
第二共同発明者の署名	日付	Second inventor's signature		Date	
		T. Yotsugi		9/21/2001	
住所	Residence Hitachinaka, Japan				
国籍	Citizenship Japan				
私書箱	Post Office Address c/o Hitachi, Ltd., Intellectual Property Group New Marunouchi Bldg. 5-1, Marunouchi 1-chome, Chiyoda-ku, Tokyo 100-8220, Japan				
第三共同発明者		3-00		Full name of third joint inventor, if any Hiroaki ISHIZAWA	
第三共同発明者の署名	日付	Third inventor's signature		Date	
		H. Ishizawa		9/26/2001	
住所	Residence Hitachinaka, Japan				
国籍	Citizenship Japan				
私書箱	Post Office Address c/o Hitachi, Ltd., Intellectual Property Group New Marunouchi Bldg. 5-1, Marunouchi 1-chome, Chiyoda-ku, Tokyo 100-8220, Japan				
第四共同発明者		Full name of fourth joint inventor, if any			
第四共同発明者の署名	日付	Fourth inventor's signature		Date	
住所	Residence				
国籍	Citizenship				
私書箱	Post Office Address				
第五共同発明者		Full name of fifth joint inventor, if any			
第五共同発明者の署名	日付	Fifth inventor's signature		Date	
住所	Residence				
国籍	Citizenship				
私書箱	Post Office Address				

(第六以降の共同発明者についても同様に記載し、署名をすること)

(Supply similar information and signature for sixth and subsequent joint inventors.)